Information to be provided to the subject before taking biological material

Sample information form

Institutional Technical and Ethical Review Board
University of Public Health, Yangon
Ministry of Health and Sports
Republic of the Union of Myanmar

The following information should be provided to the subjects before collecting biological material from them.

Type of sample, how it will be obtained, how long it will be kept and disposed

1. Detailed procedure in sequential order.
2. Degree of invasiveness.
3. In case of invasive procedures,
   a. any additional risk
   b. arrangement for treating complications that may arise during or after invasive procedure to collect specimens.
4. Protection of privacy in all physical examination.
5. How long the samples will be kept?
6. Arrangements for final disposal of the samples at the end of the research study.

Type of consent to be obtained (Explain briefly)

Fully restricted	: Consent for the specific research study only
Partially restricted consent	: Consent for this research study and future research studies related to this study
Unrestricted consent	: Consent for this research study and future research studies unrelated to this study

Whether identity will be retained or not? (Explain briefly)

1. Unidentified (anonymous or anonymized).
2. Coded (linked or identifiable).
3. Identified.

How will confidentiality be ensured?

1. How confidentiality and privacy of personal information will be protected?
2. Where samples and clinical information will be kept?
3. Who will have access to samples and research results?
4. Whether the results of research will be relayed back to the research subject?